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Report of EPO 03018636.3
Your Ref.: F030867



(11) **EP 1 281 537 A1**

(12) **EUROPEAN PATENT APPLICATION**
published in accordance with Art. 158(3) EPC

(43) Date of publication:
05.02.2003 Bulletin 2003/06

(51) Int Cl.7: **B41M 5/26, G11B 7/24**

(21) Application number: **01930074.8**

(86) International application number:
PCT/JP01/03949

(22) Date of filing: **11.05.2001**

(87) International publication number:
WO 01/085464 (15.11.2001 Gazette 2001/46)

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR**
Designated Extension States:
AL LT LV MK RO SI

- **INOUE, Hiroyasu, TDK Corporation**
Tokyo 103-8272 (JP)
- **KATO, Tatsuya, TDK Corporation**
Tokyo 103-8272 (JP)
- **UTSUNOMIYA, Hajime, TDK Corporation**
Tokyo 103-8272 (JP)
- **TANAKA, Yoshitomo, TDK Corporation**
Tokyo 103-8272 (JP)

(30) Priority: **12.05.2000 JP 2000139538**
23.06.2000 JP 2000190315

(71) Applicant: **TDK Corporation**
Chuo-ku, Tokyo 103-8272 (JP)

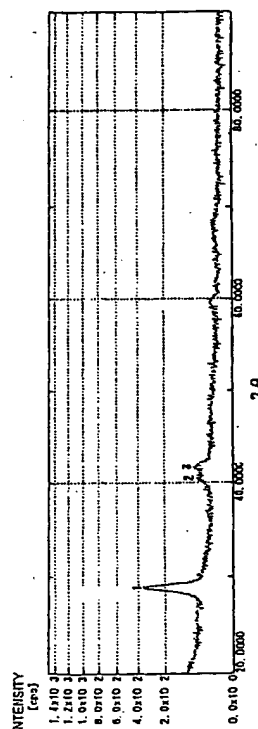
(74) Representative: **Wise, Stephen James et al**
c/o RAWORTH, MOSS & COOK,
Raworth House,
36 Sydenham Road
Croydon, Surrey CR0 2EF (GB)

(72) Inventors:
• **SHINGAI, Hiroshi, TDK Corporation**
Tokyo 103-8272 (JP)

(54) **OPTICAL RECORDING MEDIUM**

(57) The optical recording medium of the invention has a phase change recording layer containing Sb as a main component. When the recording layer is crystallized, the crystallized region contains rhombohedral crystals consisting essentially of Sb and is substantially free of a crystal phase other than the rhombohedral crystals. The optical recording medium of the invention has a high transfer rate and its recording layer is improved in thermal stability.

FIG. 6



EP 1 281 537 A1

Description

TECHNICAL FIELD

5 [0001] This invention relates to a phase change optical recording medium.

BACKGROUND ARTS

10 [0002] Great attention is now paid to optical recording media capable of high density recording and erasing the once recorded information for rewriting. Among such rewritable optical recording media, phase change recording media are designed such that recording is performed by irradiating a laser beam to a recording layer to change its crystalline state and reading is performed by detecting the change of reflectivity of the recording layer associated with that state change. The phase change recording media are of greater interest because the drive unit used for their operation may have a simple optical system as compared with that used for magneto-optical recording media.

15 [0003] For the phase change recording layer, calcogenide materials such as Ge-Sb-Te are often used because of a greater difference in reflectivity between the crystalline and amorphous states and a relatively high stability in the amorphous state.

[0004] When information is recorded in a phase change optical recording medium, the recording layer is irradiated with a laser beam having a high power (recording power) sufficient to heat the recording layer at or above its melting point. In the region where the recording power is applied, the recording layer is melted and then rapidly cooled, forming a recorded mark in the amorphous state. To erase the recorded mark, the recording layer is irradiated with a laser beam having a relatively low power (erasing power) sufficient to heat the recording layer above its crystallization temperature, but below its melting point. The recorded mark to which the erasing power is applied is heated above the crystallization temperature and then slowly cooled, resuming the crystalline state. Therefore, the phase change optical recording medium allows for overwriting simply by modulating the intensity of a single laser beam.

25 [0005] In order to increase the recording density and transfer rate of a recording medium, attempts have been made to reduce the wavelength of recording/reading beam, to increase the numerical aperture of an objective lens in a recording/reading optical system, and to increase the linear velocity of the medium. When a recording laser beam is irradiated to a medium rotating at a linear velocity V , the recording laser beam defines on the surface of the recording layer a spot having a diameter represented by λ/NA wherein λ is the wavelength of the laser beam and NA is the numerical aperture of the objective lens. The spot diameter λ/NA divided by the linear velocity V , i.e., $(\lambda/NA)/V$ gives the time of irradiation of laser beam to the recording layer, that is, the time taken for passage across a beam spot. As the recording density and transfer rate increase, the irradiation time of laser beam to the recording layer becomes shorter and shorter. This makes it difficult to optimize overwriting conditions.

35 [0006] Problems arising from overwriting at an increased linear velocity are discussed below.

[0007] An increased linear velocity leads to a shortened irradiation time of recording laser beam. It is then a common practice to increase the recording power in proportion to the increased linear velocity for preventing the heated temperature of the recording layer from lowering.

40 [0008] On the other hand, to erase the amorphous recorded mark (to recrystallize), an erasing beam must be irradiated such that the recording layer may be held for at least a predetermined time at a temperature between the crystallization temperature and the melting point. The attempt to increase the erasing power in proportion to the increased linear velocity for preventing the heated temperature of the recording layer from lowering has a less likelihood to erase the recorded mark because the irradiation time is reduced as a result of the increased linear velocity.

45 [0009] Therefore, to increase the linear velocity for improving the transfer rate, the recording layer must be formed of a composition having a relatively high crystal transition speed such that recrystallization is completed within a relatively short time (as disclosed in JP-A 1-78444 and JP-A 10-326436).

DISCLOSURE OF THE INVENTION

50 [0010] The recording layer featuring a high crystal transition speed, that is, a short time required for crystallization, however, is thermally less stable. That is, the recording layer suffers from the problem of low storage reliability since it readily crystallizes in a relatively hot environment.

[0011] Also, the high transfer rate can be established by such a method as by increasing the linear velocity of the medium or by increasing the linear recording density of the medium. The inventors found that reducing the recorded mark length in order to increase the linear recording density sacrifices the thermal stability of the recorded mark.

55 [0012] An object of the invention is to provide a phase change optical recording medium having a recording layer with an increased transfer rate and a high thermal stability.

[0013] According to the invention, there is provided an optical recording medium comprising a phase change record-

ing layer containing antimony as a main component, the recording layer being able to be crystallized to provide a crystallized region which contains rhombohedral crystals consisting essentially of antimony and is substantially free of a crystal phase other than the rhombohedral crystals.

[0014] Such objects are achievable by the following embodiments (1) to (3) of the invention.

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(1) An optical recording medium comprising a phase change recording layer containing antimony as a main component, the recording layer being able to be crystallized to provide a crystallized region which contains rhombohedral crystals consisting essentially of antimony and is substantially free of a crystal phase other than the rhombohedral crystals.

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(2) The optical recording medium of (1) wherein said recording layer further contains tellurium and/or indium as a main component.

(3) The optical recording medium of (1) or (2) wherein said recording layer contains at least one element selected from the group consisting of rare earth elements, zirconium, hafnium, titanium and tin as an auxiliary component, and

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said medium further comprises a dielectric layer disposed contiguous to said recording layer and in front of said recording layer as viewed from the side where a recording/reading beam enters, said dielectric layer containing silicon oxide, silicon nitride, aluminum oxide, or a mixture of zinc sulfide and silicon oxide, the content of silicon oxide in the mixture being at least 30 mol%.

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FUNCTION

[0015] In the phase change recording layer containing antimony (Sb) as a main constituent component, the crystal transition speed increases as the Sb content increases. On the other hand, the thermal stability of the recording layer lowers as the Sb content increases. To improve thermal stability, it is preferable to reduce the Sb content and instead, add a thermal stability-enhancing element. This, in turn, makes it difficult to increase the crystal transition speed of the recording layer.

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[0016] In the medium of the invention, the phase change recording layer provides a crystallized region which contains rhombohedral crystals consisting essentially of Sb and is substantially free of a crystal phase other than the rhombohedral crystals. If the crystalline phase included in the phase change recording layer based on Sb contains only rhombohedral crystals consisting essentially of Sb, then this recording layer provides a higher crystal transition speed than a recording layer having the same Sb content and consisting of face-centered cubic (f.c.c.) crystals and also a higher crystal transition speed than a recording layer having the same Sb content and consisting of Sb and Sb₂Te₃ phases. Accordingly, the invention is successful in increasing the crystal transition speed without extremely increasing the Sb content. Therefore, the invention entails a phase change optical recording medium capable of overwriting at a high linear velocity and having improved thermal stability.

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[0017] In one preferred embodiment, at least one element selected from among rare earth elements, Zr, Hf, Ti and Sn is added to the recording layer as an auxiliary component, thereby elevating the crystallization temperature of the recording layer. This ensures that the recorded mark is fully thermally stable even when the recording layer is of a high crystal transition speed composition, and when the recorded mark is made short. The medium is thus improved in storage reliability.

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[0018] There are known additive elements capable of elevating the crystallization temperature of a recording layer. Most additive elements, however, function to lower the crystal transition speed of the recording layer. In contrast, by adding an auxiliary component element such as rare earth element to the recording layer and at the same time, disposing a dielectric layer of a specific composition contiguous to the recording layer and in front of the recording layer as viewed from the side where a recording/reading laser beam enters the medium, the preferred embodiment of the present invention is successful in improving the thermal stability and increasing the crystal transition speed of the recording layer. The invention medium is advantageous especially in high linear velocity recording.

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[0019] It is known that rare earth elements may be added to a phase change recording layer containing at least Sb. Some of the patent references to be discussed below describe that rare earth elements are effective for improving the crystallization temperature of recording layers. The patent references below, however, do not describe the recording layer containing only rhombohedral crystals consisting essentially of Sb as the crystal phase. Nor they describe the combination of the addition of rare earth elements with the dielectric layer of the specific composition prescribed by the present invention.

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[0020] JP-A 2-3113 discloses an information recording thin film formed of In₄₄Sb₄₆Bi₁₀ (atomic ratio) by evaporation to a thickness of 100 nm. In Examples thereof, thin films based on this composition to which Nd was added were prepared and measured for phase change temperature. A rise of phase change temperature due to Nd addition was ascertained in this way.

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[0021] JP-A 2-35636 and 2-151481 disclose phase change information recording thin films containing Sb, Te and other elements. It is described that the crystallization temperature can be elevated by adding rare earth elements or analogues. In Examples thereof, however, no thin films having rare earth elements added were prepared.

[0022] JP-A 10-326436 describes that rare earth elements may be added to a phase change recording layer containing Sb and Te. It is described that Zn, Cu, Au, Ag, Pd, Pt, Cr, Co, Zr, Ti, Mn, Mo, Rh and rare earth elements themselves or compounds thereof with Sb or Te have high melting points so that they precipitate as fine dispersed clusters and serve as crystal nuclei, contributing to high speed crystallization. Examples therein lack samples having rare earth elements added. Recording at a wavelength 780 nm, a numerical aperture NA of 0.55 and a linear velocity of 4.8 m/s or below is described in Examples, which indicates that high linear velocity recording as contemplated in the present invention is out of consideration.

[0023] JP-A 2000-43415 discloses a phase change recording layer containing Sb and Te and having a metastable Sb_3Te phase belonging to the space group $\text{Fm}\bar{3}\text{m}$. This Sb_3Te phase has a face-centered cubic (f.c.c.) structure as described therein. It is also described that by adding nitrogen, boron, carbon, rare earth element or transition metal element to the recording layer, an improvement is made in the long-term storage of initially recorded data. However, Examples therein lack samples having rare earth elements added. It is also noted that in Examples, the linear velocity for recording is 7 m/s.

[0024] JP-A 2000-52657 discloses a phase change recording layer containing Sb, Te, Group Ib element and Group IIb element and having a metastable phase belonging to the space group $\text{Fm}\bar{3}\text{m}$. This metastable phase is a f.c.c. structure phase as typified by Sb_3Te phase. Although this patent describes that rare earth elements may be added to the recording layer, Examples therein lack samples having rare earth elements added. In Examples, the linear velocity for recording is 8 m/s or below. It is also described that the content of Ag+Au in the recording layer is adjusted in accordance with the linear velocity. Namely, in this patent, compositions free of Ag and/or Au are out of consideration. More specifically, Table 2 in this patent publication describes a recording layer 14 of $\text{Ag}_{0.05}\text{In}_{0.04}\text{Sb}_{0.61}\text{Te}_{0.3}$ containing a f.c.c. phase, Sb_2Te_3 phase and Sb phase. That is, this recording layer is a mixture of a f.c.c. phase and a rhombohedral crystal phase. It is described that in evaluating the recording layer 14, Sb and Sb_2Te_3 --which develop as precipitates when the metastable phase is differentiated-- precipitate in the recording layer, which is inadequate for high density recording.

[0025] JP-A 9-71049 discloses an optical information recording medium having a recording layer consisting of Sb, Te and M wherein M is at least one element of Ag, Cu and Au, wherein these constituent elements are present as $\text{Sb}_x\text{Te}_{1-x}$ wherein $0.70 < x < 0.90$ and $(\text{M}_z\text{Te}_{1-z})_b\text{Sb}$ wherein $0 < z < 0.33$ and $0 < b < 1$. It is described in paragraph 0013 of the patent publication that on X-ray analysis, a sharp peak of Sb rather than that of Sb_2Te_3 appears. That is, this recording layer contains both a Sb_2Te_3 phase and a Sb phase as crystals, but not a Sb phase alone. Therefore, the recording layer of this patent publication differs from the recording layer of the present invention and fails to achieve the desired effects of the present invention. It is not described in this patent publication that auxiliary component elements as prescribed in the present invention are added to the recording layer.

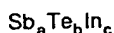
BRIEF DESCRIPTION OF THE DRAWINGS

[0026] FIGS. 1, 2, 3 and 4 are cross-sectional views of various embodiments of optical recording media. FIG. 5 is an electron beam diffraction pattern of the recording layer of an optical recording medium according to the invention. FIGS. 6 and 7 are X-ray diffraction patterns of the recording layer of the optical recording medium according to the invention.

BEST MODE FOR CARRYING OUT THE INVENTION

[0027] The optical recording medium of the invention includes a phase change recording layer which contains antimony (Sb) as a main component. Since use of Sb alone as a main component can result in too low a crystallization temperature and too low thermal stability, it is preferable to use tellurium (Te) and/or indium (In) as an additional main component. Of these, Te is especially preferable because of a possibility to increase the degree of modulation. Also preferably, the recording layer contains at least one element selected from among rare earth elements, zirconium, hafnium, titanium and tin as an auxiliary component. It is noted that the term "rare earth elements" used herein include yttrium (Y), scandium (Sc) and lanthanoids.

[0028] Where the atomic ratio of the main component constituent elements is represented by the formula I:



wherein $a+b+c = 1$, it is preferred that a range from 0.6 to 0.9, b range from 0 to 0.4, and c range from 0 to 0.4.

[0029] In formula I, too small a value of "a" representative of the Sb content may entail a greater reflectivity difference associated with phase change, but a sharp decline of crystal transition speed to impede erasure. On the other hand, with too large values of "a," the crystallization temperature may lower so that the thermal stability of recorded marks is aggravated. Also, too large a value of "a" may entail a smaller reflectivity difference associated with phase change and hence, a lower degree of modulation.

[0030] Like Sb, the auxiliary component elements are effective for increasing the crystal transition speed when a dielectric layer of specific composition is disposed in front of the recording layer as viewed from the recording/reading beam incident side. Therefore, by substituting the auxiliary component element(s) for a portion of Sb, the crystal transition speed is increased, and at the same time, the thermal stability of recorded marks is improved. Therefore, the invention medium performs well particularly when operated at a high linear velocity.

[0031] The content of the auxiliary component element in the recording layer is preferably up to 30 at%, more preferably up to 25 at%. Too high an auxiliary component element content may lead to too high a crystallization temperature, which hinders to initialize or crystallize an amorphous recording layer immediately after its formation. In order that the auxiliary component element added fully exert the effects of increasing the crystal transition speed and the recorded mark's thermal stability, the content of auxiliary component element should preferably be set at 0.1 at% or above, more preferably 0.5 at% or above.

[0032] The preferred auxiliary component elements are rare earth elements since they are more effective for increasing the crystal transition speed and the recorded mark's thermal stability.

[0033] In addition to the above-mentioned main and auxiliary components, the recording layer may contain one or more other element if desired. Such an additive element is designated element M wherein M is at least one element selected from among Ag, Au, Bi, Se, Al, P, Ge, H, Si, C, V, W, Ta, Zn, Pb and Pd. Element M is added, if necessary, for achieving any desired effect. For example, at least one of V and Ta is preferably added for the purpose of improving durability against rewriting, more specifically for suppressing any loss of erasability by repetitive rewriting. Also, Ge is preferably added for the purpose of improving thermal stability.

[0034] For the purpose of increasing the degree of modulation, it is recommended that the addition of element M be avoided. Even when element M is added, the content of element M in the recording layer should preferably be suppressed to 10 at% or less. Too high an element M content may lead to a small change of reflectivity associated with phase change, failing to provide a degree of modulation.

[0035] Preferably the recording layer has a thickness of 2 to 50 nm, more preferably 2 to 30 nm. Too thin a recording layer may impede the growth of a crystal phase, resulting in an insufficient change of reflectivity associated with phase change. A too thick recording layer possesses a large heat capacity which may impede recording, and has a low reflectivity and a low degree of modulation.

[0036] The composition of the recording layer can be analyzed by electron probe microanalysis (EPMA), x-ray microanalysis and inductively coupled plasma emission spectroscopy (ICP), for example.

[0037] The recording layer is preferably formed by a sputtering process. The sputtering conditions are not critical. When a material containing plural elements is to be deposited by sputtering, an alloy target may be used. A multi-source sputtering process using a plurality of targets is also useful.

[0038] The recording layer contains rhombohedral crystals consisting essentially of antimony (Sb). For increasing the crystal transition speed of the recording layer, the invention employs a relatively high Sb content and substantially excludes from the recording layer a crystal phase other than the rhombohedral crystals. The rhombohedral crystals consisting essentially of Sb are present as a phase consisting solely of Sb and an Sb phase forming a solid solution with another element. It is required that at least one of these phases be present in the recording layer. It is noted that by the term "rhombohedral crystals consisting essentially of Sb," a Sb_2Te_3 phase is not encompassed.

[0039] The presence of rhombohedral crystals consisting essentially of Sb and the substantial absence of another crystal phase can be ascertained by electron beam diffraction and x-ray diffraction. The recording layer substantially free of a crystal phase other than the rhombohedral crystals, as used herein, means that on electron beam diffraction and x-ray diffraction analysis, the crystal phase other than the rhombohedral crystals is not detected.

[0040] As long as the above-described composition and crystal phase requirements are met, other factors of the recording layer are not critical. The optical recording medium may have any desired structure as long as it satisfies the composition and crystal phase requirements.

[0041] One general construction of the phase change optical recording medium is illustrated in FIG. 1 as comprising a substrate 2, and a first dielectric layer 31, a recording layer 4, a second dielectric layer 32, a reflective layer 5, and a protective layer 6 stacked successively on the substrate 2 in the described order. In this medium, a recording/reading beam is irradiated through the substrate 2.

[0042] Also, the optical recording medium may be constructed as shown in FIG. 2, such that a recording/reading beam is irradiated to the recording layer without passing through the substrate 2. In this embodiment, a reflective layer 5, a second dielectric layer 32, a recording layer 4, a first dielectric layer 31 are stacked on a substrate 2 in the described order, and a protective layer 6 of a light-transmitting material such as resin is finally laid thereon. A recording/reading

beam is irradiated to the recording layer 4 through the protective layer 6.

[0043] In one preferred embodiment, an auxiliary component element such as rare earth element is added to the recording layer. In order that the auxiliary component element added function to increase the crystal transition speed, the dielectric layer disposed in front of the recording layer as viewed from the recording/reading beam incident side should have a specific composition. This specific composition is a composition containing silicon oxide, silicon nitride, aluminum oxide, or a mixture of zinc sulfide and silicon oxide. The composition should preferably contain at least 30 mol%, more preferably at least 40 mol%, and most preferably 100 mol% of any of these compounds. That is, the most preferred composition is silicon oxide, silicon nitride, aluminum oxide, or a mixture of zinc sulfide and silicon oxide. In the mixture, the content of silicon oxide should be at least 30 mol%, preferably 30 to 70 mol% of the zinc sulfide and silicon oxide combined. If the first dielectric layer disposed in front of the recording layer is outside the specific composition, the crystal transition speed-up effect of auxiliary component element addition is not exerted.

[0044] The dielectric layer of the specific composition may be the entirety of the first dielectric layer 31 in FIGS. 1 and 2. Where the first dielectric layer 31 is a laminate of dielectric sublayers, it suffices that at least the dielectric sublayer disposed contiguous to the recording layer 4 has the specific composition. Such an embodiment is illustrated in FIG. 3. The first dielectric layer 31 in the medium shown in FIG. 3 is a laminate of two dielectric sublayers 311 and 312. The medium has the same structure as the medium of FIG. 2 except for the first dielectric layer 31.

[0045] The thickness of the first and second dielectric layers 31 and 32 may be determined as appropriate to achieve the effect of protecting the recording layer 4 and the effect of improving the degree of modulation. Usually, the first dielectric layer 31 preferably has a thickness of 30 to 300 nm, more preferably 50 to 250 nm; and the second dielectric layer 32 preferably has a thickness of 5 to 50 nm. In order that the auxiliary component element added function to fully increase the crystal transition speed, the dielectric layer of the specific composition should preferably have a thickness of at least 1.5 nm, with a thickness of 2.5 nm or more achieving the best results. Since the provision of such a thin dielectric layer achieves satisfactory effects, it is not required that the first dielectric layer 31 in its entirety have the specific composition. Then the provision of a dielectric layer of the specific composition does not affect the freedom of optical and thermal design of the first dielectric layer 31 as a whole.

EXAMPLE

[0046] Examples of the invention are given below by way of illustration.

Example 1

[0047] An optical recording disk sample of the structure illustrated in FIG. 3 was prepared by injection molding polycarbonate into a disk-shaped substrate 2 having a diameter of 120 mm and a thickness of 1.1 mm in which grooves were formed simultaneous with injection molding. On the surface of the substrate 2, a reflective layer 5, a second dielectric layer 32, a recording layer 4, a first dielectric layer 31 and a protective layer 6 were successively formed by the following procedure.

[0048] The reflective layer 5 was formed by sputtering in an argon atmosphere. The target used was $\text{Ag}_{98}\text{Pd}_1\text{Cu}_1$ (atomic ratio). The reflective layer was 100 nm thick.

[0049] The second dielectric layer 32 was formed by sputtering a target of Al_2O_3 in an argon atmosphere. The second dielectric layer was 20 nm thick.

[0050] The recording layer 4 was formed by sputtering in an argon atmosphere. It was 12 nm thick. The composition of the recording layer is shown in Table 1.

[0051] As the first dielectric layer 31, a dielectric sublayer 311 of 50 mol% ZnS-50 mol% SiO_2 and 5 nm thick was first formed in close contact with the recording layer 4. Another dielectric sublayer 312 of 80 mol% ZnS-20 mol% SiO_2 and 120 nm thick was then formed on the dielectric sublayer 311. These sublayers were formed by sputtering in an argon atmosphere.

[0052] The protective layer 6 was formed by joining a polycarbonate sheet of 100 μm thick with a UV-curable adhesive.

[0053] The recording layers of the disk samples were initialized or crystallized by means of a bulk eraser. Each disk sample was mounted on an optical recording medium tester where recording was carried out under the following conditions.

laser wavelength: 405 nm,

numerical aperture NA: 0.85,

recording linear velocity: 11.4 m/s

recording signals: single signals having a frequency corresponding to a recorded mark length 692 nm

[0054] Next, while the disk sample was rotated so as to provide a linear velocity as shown in Table 1, a laser beam at an erasing power level was irradiated thereto. A erasability was determined, with the results shown in Table 1. This

measurement was repeatedly carried out while changing the erasing power level. The maximum erasability is reported in Table 1. An erasability of less than 25 dB is regarded to be unerased.

[0055] For the evaluation of thermal stability, samples were prepared by the same procedure as the optical recording disk samples except that slide glass was used as the substrate. These samples were rested on a heating stage. While the sample was heated at 30°C/min, light is irradiated to the recording layer through the substrate. The temperature at which the reflectivity changed was determined and reported as the crystallization temperature of the recording layer. The results are shown in Table 1.

[0056] By dissolving away the substrate and the layers other than the recording layer from the optical recording disk samples, there were left only the recording layers. The recording layers were examined for crystal phase by performing electron beam diffraction under a transmission electron microscope. As a result, for all the samples, rhombohedral crystals consisting essentially of Sb were detected, but not any crystal phase (e.g., Sb_2Te_3 phase) other than the rhombohedral crystals. FIG. 5 shows the electron beam diffraction pattern of sample No. 4 by this type of analysis.

[0057] The recording layers were also subjected to in-plane diffraction analysis using a thin film x-ray diffractometer ATX-G by Rigaku Denki Co., Ltd. As a result, for all the samples, rhombohedral crystals consisting essentially of Sb were detected, but not any crystal phase (e.g., Sb_2Te_3 phase) other than the rhombohedral crystals. FIG. 6 shows the x-ray diffraction pattern of sample No. 4 by this type of analysis.

[0058] Separately, samples for x-ray diffraction were prepared by forming only the recording layers on the surface of the substrates 2. The recording layers were formed by the same procedure as the above optical recording disk samples except that the thickness was 200 nm. These samples were analyzed by x-ray diffraction for examining crystal phases. The crystal phase detected herein was just the same as in the above electron beam diffraction and in-plane diffraction. FIG. 7 shows the x-ray diffraction pattern of sample No. 4 by this type of analysis.

[0059] It is noted that in the patterns of FIGS. 6 and 7, all the peaks are attributable to the Sb phase. Judging from the position of each peak, it is believed that the elements other than Sb form a solid solution with the Sb phase. In the patterns of FIGS. 6 and 7, the peaks designated at 1, 2 and 3 correspond to (012), (104) and (110) planes, respectively.

Table 1

Sample No.	Recording layer composition (at%)							Erasability (dB)			Crystallization temperature (°C)
	Sb	Te	In	Ge	Tb	Dy	Gd	Linear velocity	Linear velocity	Linear velocity	
1	69.6	30.4	—	—	—	—	—	33.8	13.8	4.6	163
2	67.3	30.3	—	—	2.4	—	—	37.6	26.9	8.4	184
3	65.6	30.4	—	—	4.0	—	—	37.0	33.7	28.3	230
4	76.0	16.9	1.1	6.0	—	—	—	33.6	21.8	13.8	206
5	73.0	16.1	1.1	5.8	4.0	—	—	unmeasurable	36.5	28.0	250
6	72.9	16.1	1.1	5.8	—	4.1	—	unmeasurable	30.1	30.9	237
7	73.0	16.1	1.1	5.8	—	—	4.0	unmeasurable	35.0	27.6	226

[0060] It is seen from Table 1 that the recording layers containing rhombohedral crystals as the crystal phase achieve a satisfactory erasability even at a high linear velocity of 11.4 m/s or higher. The recording layer of sample No. 4 having Ge added as the auxiliary component shows an elevated crystallization temperature. The recording layers having a rare earth element added as the auxiliary component achieves improvements in both erasable linear velocity and thermal stability.

[0061] It is noted in Table 1 that "unmeasurable" under the heading of erasability indicates that recording is impossible because the crystal transition speed of the recording layer is extremely high.

Example 2

Sample No. 8

[0062] An optical recording disk, sample No. 8, of the structure illustrated in FIG. 4 was prepared by the following procedure. The structure of FIG. 4 is the same as that of FIG. 3 except that the first dielectric layer 31 includes three dielectric sublayers 311, 312 and 313.

[0063] The substrate 2 and the reflective layer 5 were the same as in Example 1.

[0064] The second dielectric layer 32 was formed by reactive sputtering in a mixed atmosphere of Ar and N₂ using an aluminum target. The second dielectric layer was 20 nm thick.

[0065] The recording layer 4 was formed by sputtering in an argon atmosphere. It was 12 nm thick. The composition of the recording layer was Sb_{72.5}Te_{13.7}In_{0.9}Ge_{4.9}Tb₈ (at%).

[0066] The dielectric sublayer 311 in close contact with the recording layer 4 was formed of Al₂O₃ to a thickness of 5 nm. The dielectric sublayer 312 was formed of 80 mol% ZnS-20 mol% SiO₂ to a thickness of 63 nm. The dielectric sublayer 313 was formed of aluminum nitride to a thickness of 100 nm.

[0067] The protective layer 6 was formed by joining a polycarbonate sheet of 100 μm thick to the dielectric layer 31 with a UV-curable adhesive.

Sample No. 9

[0068] An optical recording disk sample No. 9 was prepared by the same procedure as sample No. 8 except that the first dielectric layer 31 had a two-layer structure as shown in FIG. 3. The dielectric sublayer 311 in contact with the recording layer 4 was formed of 80 mol% ZnS-20 mol% SiO₂ to a thickness of 68 nm, and the dielectric sublayer 312 was formed of aluminum nitride to a thickness of 100 nm.

Evaluation

[0069] The samples prepared above were measured for erasability as in Example 1. The results are shown in Table 2.

Table 2

Sample No.	Dielectric sublayer 311 composition	Erasability (dB)	
		Linear velocity	
		22.8m/s	27.6m/s
8	Al ₂ O ₃	26.5	18.4
9	ZnS(80)-SiO ₂ (20)	8.5	5.8

[0070] It is seen from Table 2 that sample No. 9 in which the dielectric sublayer 311 in contact with the recording layer 4 was formed of 80 mol% ZnS-20 mol% SiO₂ has a very low erasability whereas sample No. 8 in which the dielectric sublayer 311 is formed of Al₂O₃ has a high erasability. These results demonstrate the effects resulting from the addition of rare earth element Tb to the recording layer 4 and the construction of the dielectric sublayer 311 from the specific composition.

[0071] It is noted that on analysis by electron beam diffraction and x-ray diffraction of the recording layer 4 of sample Nos. 8 and 9 as in Example 1, no crystal phase other than the rhombohedral crystals consisting essentially of Sb was detected.

[0072] Reasonable modifications and variations are possible from the foregoing disclosure without departing from either the spirit or scope of the present invention as defined by the claims.

INDUSTRIAL APPLICABILITY OF THE INVENTION

[0073] According to the invention, a phase change optical recording medium capable of overwriting at a high linear velocity and having improved thermal stability is obtainable.

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Claims

- 10 1. An optical recording medium comprising a phase change recording layer containing antimony as a main component, the recording layer being able to be crystallized to provide a crystallized region which contains rhombohedral crystals consisting essentially of antimony and is substantially free of a crystal phase other than the rhombohedral crystals.
- 15 2. The optical recording medium of claim 1 wherein said recording layer further contains tellurium and/or indium as a main component.
- 20 3. The optical recording medium of claim 1 or 2 wherein said recording layer contains at least one element selected from the group consisting of rare earth elements, zirconium, hafnium, titanium and tin as an auxiliary component, and
said medium further comprises a dielectric layer disposed contiguous to said recording layer and in front of said recording layer as viewed from the side where a recording/reading beam enters, said dielectric layer containing silicon oxide, silicon nitride, aluminum oxide, or a mixture of zinc sulfide and silicon oxide, the content of silicon oxide in the mixture being at least 30 mol%.

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FIG. 1

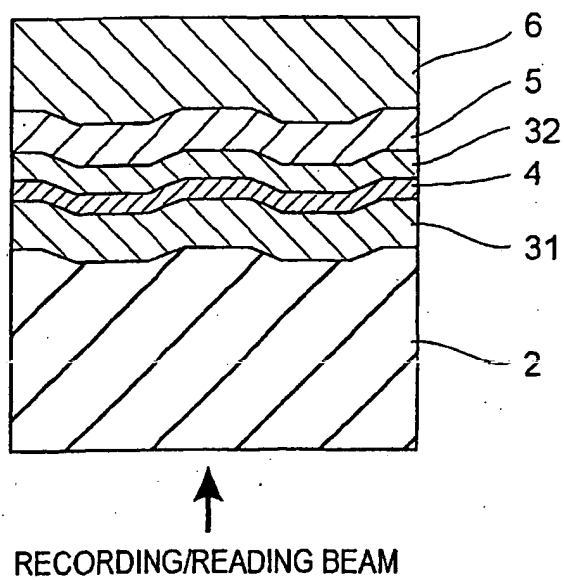


FIG. 2

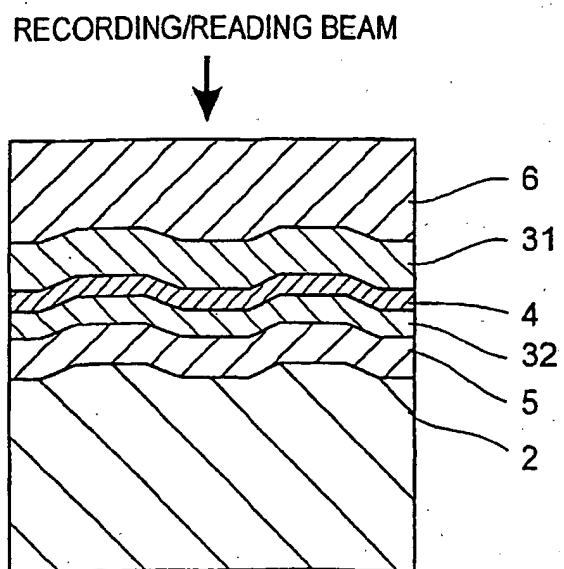


FIG. 3

RECORDING/READING BEAM

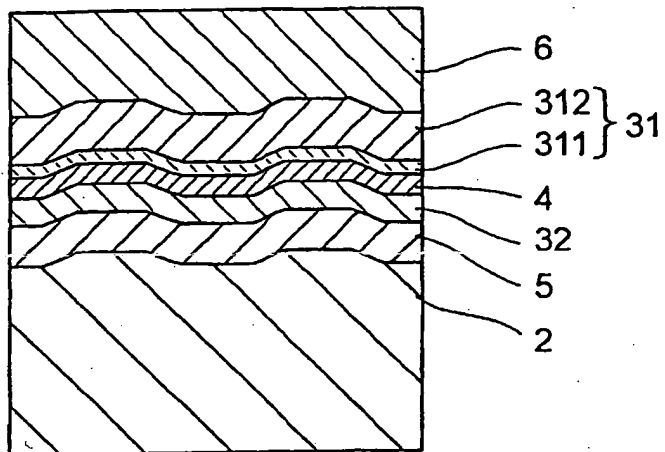


FIG. 4

RECORDING/READING BEAM

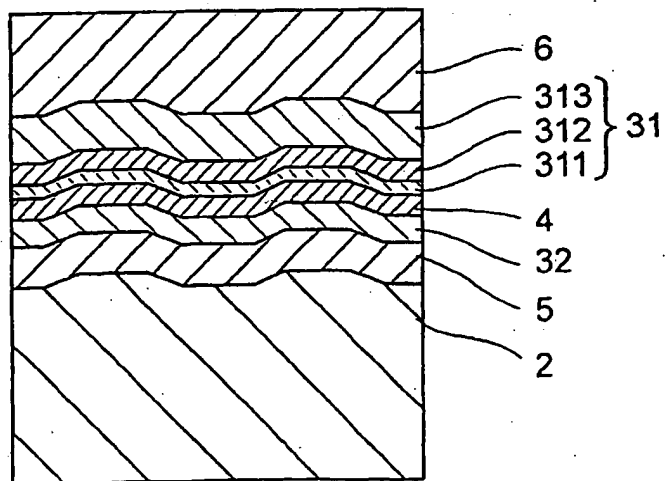


FIG. 5

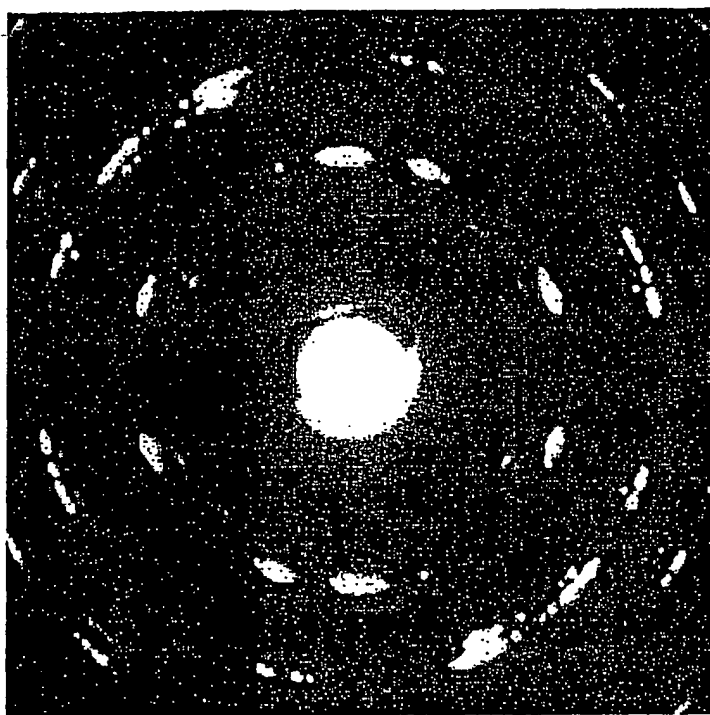


FIG. 6

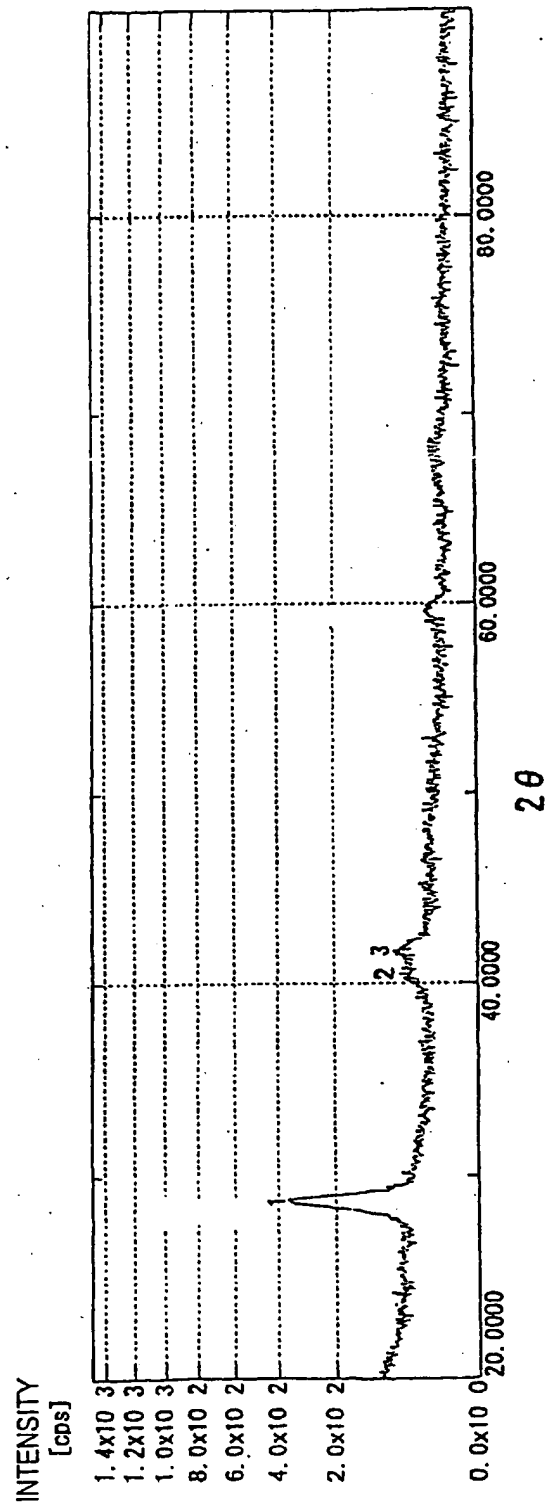
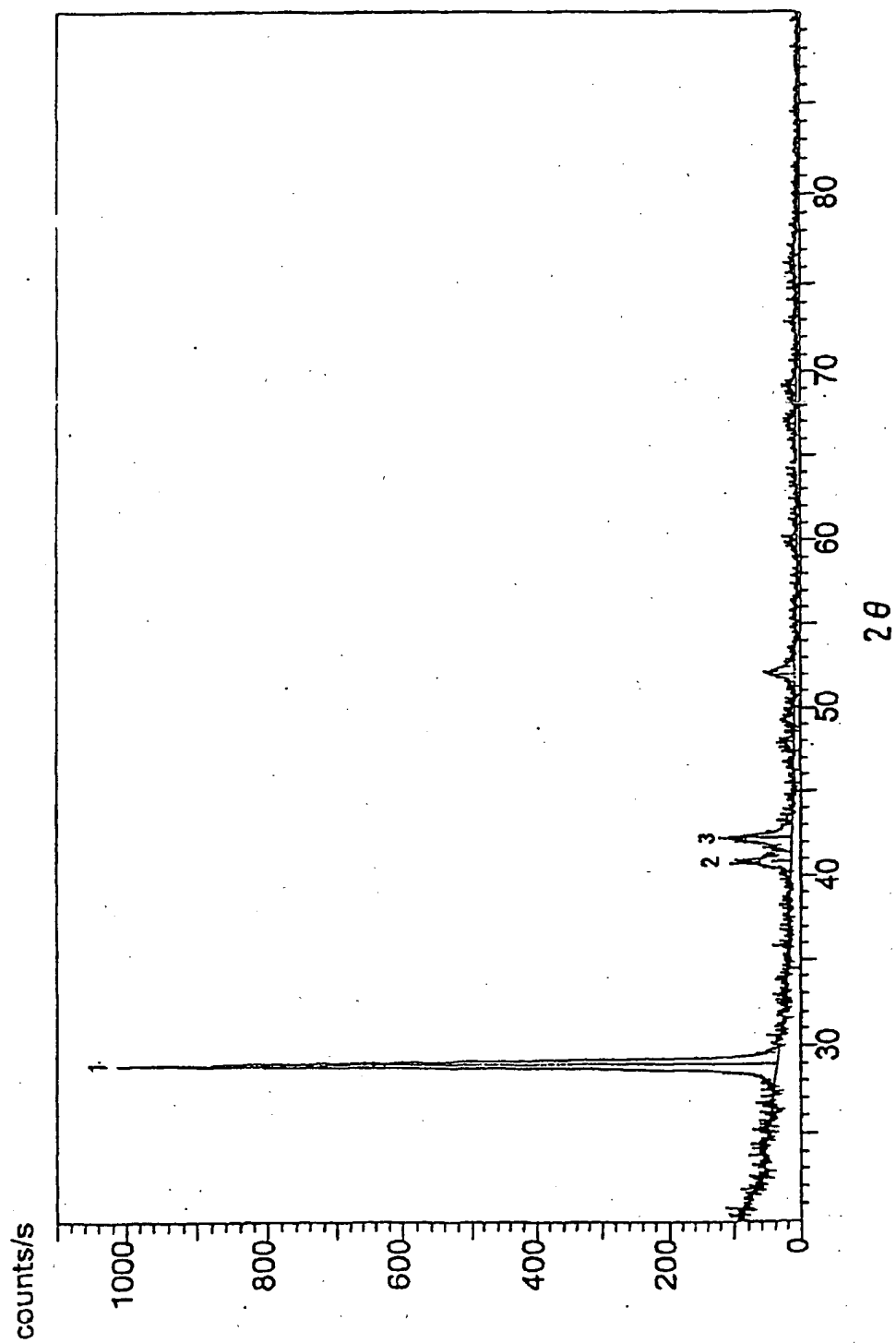


FIG. 7



INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP01/03949

A. CLASSIFICATION OF SUBJECT MATTER
Int.Cl.¹ B41M5/26, G11B7/24

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
Int.Cl.¹ B41M5/26, G11B7/24

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
Jitsuyo Shinan Koho 1922-1996 Toroku Jitsuyo Shinan Koho 1994-2001
Kokai Jitsuyo Shinan Koho 1971-2001 Jitsuyo Shinan Toroku Koho 1996-2001

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X Y	JP 2000-43415 A (Ricoh Company, Ltd.), 15 February, 2000 (15.02.00), Full text; all drawings & EP 957476 A1	1, 2 3
X Y	JP 9-71049 A (Mitsubishi Chemical Corporation), 18 March, 1997 (18.03.97), Claims; Par. Nos. [0010] to [0018], [0028] to [0043] (Family: none)	1, 2 3
X Y	JP 64-32438 A (Nippon Columbia Co., Ltd.), 02 February, 1989 (02.02.89), Full text; all drawings (Family: none)	1, 2 3
Y	JP 9-286175 A (Mitsubishi Chemical Corporation), 04 November, 1997 (04.11.97), Full text; all drawings (Family: none)	1-3

☒ Further documents are listed in the continuation of Box C.☐ See patent family annex.

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"&" document member of the same patent family

Date of the actual completion of the international search
18 July, 2001 (18.07.01)

Date of mailing of the international search report
31 July, 2001 (31.07.01)

Name and mailing address of the ISA/
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INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP01/03949

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	JP 4-316888 A (Matsushita Electric Ind. Co., Ltd.), 09 November, 1992 (09.11.92), Full text; all drawings (Family: none)	1-3

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